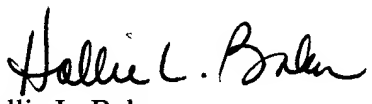
REMARKS

This amendment does not introduce new subject matter as support is found in the application as filed.

A copy of the change of the paragraph on pages 62 and 63 is attached hereto.

No fees are believed to be due with the filing of this Sequence Listing. The Commissioner is hereby authorized to debit Deposit Account No. 08-0219 any required fee necessary to maintain the pendency of this application.

Respectfully submitted,



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Single-strand rescue was conducted on the phagemid library using helper phage VCM13 and protocols provided by the supplier (Stratagene). Rescued single-stranded DNA was purified according to the method of Soares et al (Soares et al ibid) incorporating particular steps to eliminate contaminating double-stranded DNA. Controlled primer extension was conducted on purified single-stranded circular DNA using primer (SEQ ID NO.: 61) 5'ggaaacagctatgaccatg, using the conditions of Soares et al. DNA polymerase was from Boehringer, nucleotide triphosphates were from Life Technologies (Paisley, UK). a ^{32}P dCTP used as tracer was from Amersham International (Amersham, UK). Hydroxyapatite column chromatography was conducted at 60°C as described in Soares et al. Following recovery of the normalized single-stranded circles, the purified DNA was directly transformed into competent XL1-blue E.*coli* cells and plated on LB-ampicillin plates as previously described.

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